



MAY 18 2006 Doc Code: AP.PRE.REQ

PTO/SB/33 (07-05)

Approved for use through xx/xx/200x. OMB 0651-00xx

U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE

Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it displays a valid OMB control number.

PRE-APPEAL BRIEF REQUEST FOR REVIEW

Docket Number (Optional)

UTSD:798US

I hereby certify that this correspondence is being deposited with the United States Postal Service with sufficient postage as first class mail in an envelope addressed to "Mail Stop AF, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450" [37 CFR 1.8(a)]

on May 12, 2006

Signature

Typed or printed name Mark T. Garrett

Application Number

09/905,670

Filed

July 13, 2001

First Named Inventor

Phillip D. Purdy

Art Unit

3763

Examiner

Mark K. Han

Applicant requests review of the final rejection in the above-identified application. No amendments are being filed with this request.

This request is being filed with a notice of appeal.

The review is requested for the reason(s) stated on the attached sheet(s).

Note: No more than five (5) pages may be provided.

I am the

applicant/inventor.

assignee of record of the entire interest.
See 37 CFR 3.71. Statement under 37 CFR 3.73(b) is enclosed.
(Form PTO/SB/96)

attorney or agent of record.
Registration number 44,699.

attorney or agent acting under 37 CFR 1.34.

Registration number if acting under 37 CFR 1.34 _____

Signature

Mark T. Garrett

Typed or printed name

(512) 536-3031

Telephone number

May 12, 2006

Date

NOTE: Signatures of all the inventors or assignees of record of the entire interest or their representative(s) are required.
Submit multiple forms if more than one signature is required, see below*.



*Total of _____ forms are submitted.

This collection of information is required by 35 U.S.C. 132. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.11, 1.14 and 41.6. This collection is estimated to take 12 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Mail Stop AF, Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.



Arguments in Support of Pre-Appeal Brief Request for Review for 09/905,670

I. The Anticipation Rejection of Independent Claims 1 and 68 and Their Rejected Dependent Claims over Barbut Has No Factual Support

A. Claim 1

Independent claim 1 is directed to a method of navigating a spinal subarchnoid space in a living being. It comprises:

percutaneously introducing a guidewire into the spinal subarachnoid space at an entry location;

percutaneously introducing a device over the guidewire and into the spinal subarachnoid space, the device having a first passageway sized to slidably receive, and work with, at least the guidewire, and the guidewire being positioned in the first passageway; and

advancing the device over the guidewire and within the spinal subarachnoid space at least more than 10 centimeters from the entry location.

The Office asserts that this claim is anticipated by Barbut (U.S. 6,379,331) and states that the claimed guidewire is met by needle 66 disclosed in Barbut.

Barbut's needle is not a guidewire by any reasonable standard of claim interpretation. Furthermore, the Office provides no evidence that one of ordinary skill in the art—giving the term guidewire its broadest reasonable interpretation consistent with Applicant's specification—would read the term guidewire broadly enough to cover Barbut's needle. Applicant used the terms “guidewire” and “needle” differently in the application. Specification at page 15, lines 4-27.

For these reasons, the Office has failed to establish a *prima facie* case of anticipation and should withdraw the rejection of claims 1, 4-7, 11-13, 24 and 27.

B. Claim 68

Claim 68 is directed to a method of navigating a spinal subarachnoid space in a living being. It comprises:

percutaneously introducing a device into the spinal subarachnoid space at an entry location, the device having a first passageway sized to slidably receive, and work with, at least a guidewire;

advancing the device within the spinal subarachnoid space at least more than 10 centimeters from the entry location; and

accessing at least one ventricle located within the head with a second device introduced through the first passageway of the device.

The Office fails to address this claim in its rejection on page 3 of the April 20, 2006 Action, or on pages 6 or 7 of the Action where it addresses Applicant's remarks from the previous response. For this reason alone, the Office has not established a *prima facie* case of anticipation and should withdraw the rejection of claims 68 and 69. Moreover, Applicant explained in section J of the December 22, 2005 Response (pages 13 and 14) why claims 68 and 69 are patentable over Barbut.

II. The Obviousness Rejections of the Dependent Claims Have No Factual Support

A. Claim 2

As explained in Section C of the December 22, 2005 Response, the secondary reference, Michaeli, fails to cure the shortcomings of Barbut and that there is no motivation for the asserted combination. The rejection should be withdrawn.

B. Claim 3

As explained in Section D of the December 22, 2005 Response, the secondary reference, Janese, fails to cure the shortcomings of Barbut. The rejection should be withdrawn.

C. Claims 8, 19 and 20

Applicant directs the panel to its explanation in Section E of the December 22, 2005 Response of why these claims are patentable over the asserted combination. The rejection should be withdrawn.

D. Claims 17 and 22

Applicant directs the panel to its explanation in Section F of the December 22, 2005 Response of why these claims are patentable over the asserted combination. The rejection should be withdrawn.

E. Claim 25

Applicant directs the panel to its explanation in Section G of the December 22, 2005 Response of why this claim is patentable over the asserted combination. The rejection should be withdrawn.

F. Claim 28

Applicant directs the panel to its explanation in Section H of the December 22, 2005 Response of why this claim is patentable over the asserted combination. The rejection should be withdrawn.